

## (12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property  
Organization  
International Bureau



(43) International Publication Date  
22 September 2005 (22.09.2005)

PCT

(10) International Publication Number  
WO 2005/087307 A2

(51) International Patent Classification<sup>7</sup>: A61N 1/00

(21) International Application Number: PCT/IL2005/000135

(22) International Filing Date: 4 February 2005 (04.02.2005)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:

60/542,022	5 February 2004 (05.02.2004)	US
60/566,078	29 April 2004 (29.04.2004)	US

(71) Applicant (for all designated States except US): REABILITY INC. [GB/GB]; Nerine Chambers, P.O. Box 905, Road Town, Tortola (VG).

(72) Inventors; and

(75) Inventors/Applicants (for US only): EINAV, Omer [IL/IL]; 42875 Kfar Monash (IL). KORENMAN, Ernesto [IL/IL]; 7 Hativat Givati Street, 43338 Raanana (IL).

(74) Agents: FENSTER, Paul et al.; FENSTER & COMPANY, INTELLECTUAL PROPERTY LTD., P.O. Box 10256, 49002 Petach Tikva (IL).

(81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.

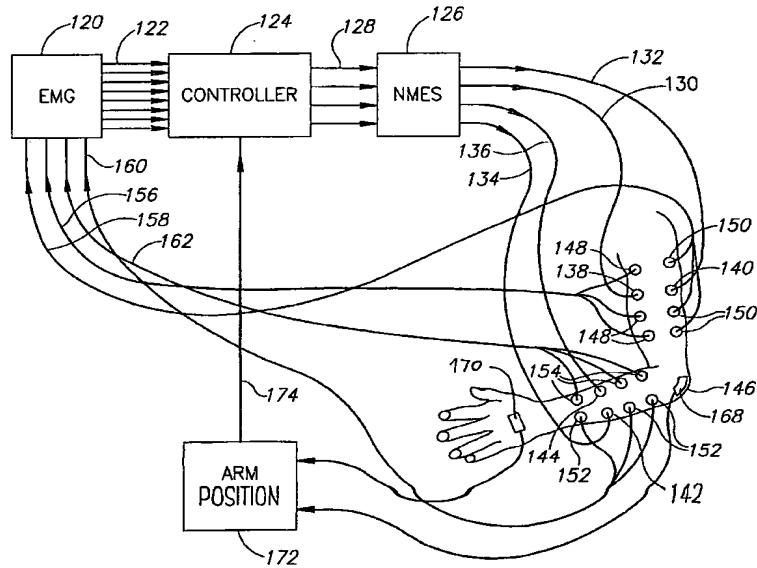
(84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

## Published:

— without international search report and to be republished upon receipt of that report

[Continued on next page]

(54) Title: NEUROMUSCULAR STIMULATION



WO 2005/087307 A2

(57) Abstract: Apparatus for rehabilitating a patient who has a paretic body part, the apparatus comprising: a) at least one electromyography (EMG) sensor adapted to being applied to a voluntary muscle of a healthy body part of the same type as the paretic body part, which at least one sensor produces at least one EMG signal; b) a neuromuscular electrical stimulation (NMES) device adapted for stimulating at least one voluntary muscle of the paretic body part; and c) a controller which controls the NMES device, making the amplitude of stimulation of the paretic body part at least partly dependent on the EMG signal from the healthy body part.

WO 2005/087307 A2



*For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.*

## NEUROMUSCULAR STIMULATION

RELATED APPLICATIONS

This application claims the benefit under 119(e) of U.S. Provisional Application No. 60/542,022, filed on February 5, 2004, and U.S. Provisional Application 60/566,078 filed on 5 April 29, 2004, the disclosure of both applications are incorporated herein by reference. This application is also related to PCT applications, being filed on same date and by the same applicant as the present application, entitled "Gait Rehabilitation Methods and Apparatuses"; "Rehabilitation with Music"; "Fine Motor Control Rehabilitation"; "Methods and Apparatuses for Rehabilitation Exercise and Training"; "Methods and Apparatus for Rehabilitation and 10 Training"; "Methods and Apparatus for Rehabilitation and Training"; "Methods and Apparatus for Rehabilitation and Training"; and having attorney docket numbers, 414/04391; 414/04396; 414/04401; 414/04388; 414/04213, 414/04404; and 414/04405, respectively. The disclosures of all these applications are incorporated herein by reference.

FIELD OF THE INVENTION

15 The field of the invention is devices for rehabilitation of patients with motor control problems.

BACKGROUND OF THE INVENTION

Voluntary muscle movement is caused by electrical impulses which originate in the somato-motor cortex of the brain. A neuron in the somato-motor cortex sends electrical signals 20 to a motor neuron in the spinal cord, which in turn sends electrical signals which stimulate the contraction of muscle fibers, producing movement. All of the muscle fibers which are stimulated by a given motor neuron are called a "motor unit." Each muscle fiber exhibits an electrical potential across its cell membrane, which changes when the muscle contracts.

In electromyography (EMG), the difference in potential on the surface of the skin is 25 measured between the center and the ends of a muscle, which gives a measure of the number of contracting muscle fibers. EMG is regularly used to diagnose a variety of medical conditions in patients, as well as in healthy subjects for research on muscle function.

In stroke patients with damage to their somato-motor cortex, electrical signals are not generated for one or more muscles or parts of muscles, or do not reach those muscles, and 30 normal contraction of those muscles is impossible. Often, residual EMG signals, too weak or too spread out to cause the muscles to contract, are still detectable.

Neuromuscular electrical stimulation (NMES) is used to produce contraction of a muscle which cannot contract normally in a stroke patient. NMES may stop spasticity in a muscle, and may prevent the muscle from atrophying. It is also known to turn NMES of a

single muscle on or off in response to residual EMG signals detected from that muscle, thereby allowing the muscle to contract under the control of the patient.

#### SUMMARY OF THE INVENTION

An aspect of an embodiment of the invention concerns applying NMES to a paretic 5 arm, or any other part of the body with voluntary muscles, in a manner which is insufficient to cause a desired motion. In an exemplary embodiment of the invention, the desired motion is provided or assisted by an actuator which moves or helps move the paretic portion. Alternatively or additionally, a patient provides the required additional neural signals (e.g., via natural pathways). In some embodiments the actuating device is used to resist motion and/or to 10 guide the motion to a desired pathway. Optionally, a controller is provided storing thereon a plurality of desired motions and expected responses by the patient to various stimulation and assistance levels.

In an exemplary embodiment of the invention, the NMES is provided at an amplitude that would be too low to produce motion by itself, but which, in combination with nerve 15 impulses arising in the patient's motor cortex, allow the arm or other body part to move, or to move more effectively than without the NMES. It does this, for example, by producing muscular feedback which helps to train the motor cortex to move that body part. In some embodiments, the NMES need not be very strong, or very precisely directed, in order to do this. Whenever this application refers to arms, it should be understood that any other body part, 20 or combination of body parts, with voluntary muscles may be used instead. Optionally, EMG signals from the same arm, or from the corresponding muscles in the patient's other arm, or from the arm of another person, are used to determine the pattern (e.g., timing and/or amplitude) of the NMES.

Another aspect of an embodiment of the invention concerns the use of EMG signals 25 from one arm, optionally a healthy arm, undergoing voluntary motion, to determine a pattern of NMES to be applied to another, paretic arm. Optionally EMG signals from the paretic arm are also used, at least to determine the timing of the NMES. Optionally, the healthy arm is the other arm of the patient, and the patient tries to move both arms in synchrony, in a mirror symmetric pattern.

30 Optionally, for either of these embodiments of the invention, the EMG and NMES involve a coordinated sequence of contractions of more than one muscle, and/or a range of amplitudes for the NMES, rather than having the NMES either on or off for a single muscle.

By providing feedback, through the kinesthetic sense, of a coordinated sequence of muscle contractions, the patient's nervous system may be encouraged to utilize alternative

undamaged pathways for nerve impulses, or alternative locations in the motor cortex, and the patient can learn to move his arm more effectively on his own. This may be especially true if the NMES is coordinated with the weak nerve impulses that the patient produces on his own, as measured by the EMG.

5        Optionally, a device, for example a robotic arm, which monitors and displays the movement of the arm, is used for the paretic arm, and optionally also for the healthy arm if one is used. Information about the movement of the arm can provide further feedback to the patient, as well as feedback for controlling the NMES, and feedback to a physical therapist who is monitoring the progress of the patient's rehabilitation. A robotic arm or similar device  
10      can also mechanically move the paretic arm, complementing the NMES by providing a different kind of kinesthetic feedback. A robotic arm can also exert a force working against the muscle, providing a way to strengthen the arm as well as to measure progress in strengthening the arm.

15      There is thus provided in accordance with an exemplary embodiment of the invention, apparatus for rehabilitating a patient who has a paretic body part, the apparatus comprising:

- a) at least one electromyography (EMG) sensor adapted to being applied to a voluntary muscle of a healthy body part of the same type as the paretic body part, which at least one sensor produces at least one EMG signal;
- b) a neuromuscular electrical stimulation (NMES) device adapted for stimulating at least one voluntary muscle of the paretic body part;
- c) a controller which controls the NMES device and expects a motion of the paretic body part, making the amplitude of stimulation of the paretic body part at least partly dependent on the EMG signal from the healthy body part such that said NMES stimulation is not sufficient, on its own, to move said paretic body part said expected motion.

25      Optionally, the at least one muscle of the healthy body part corresponds to the at least one muscle of the paretic body part.

30      In an exemplary embodiment of the invention, said controller is configured to process said EMG signals and determine at least one property of said NMES signal.

35      In an exemplary embodiment of the invention, the controller is configured so that the NMES stimulates the paretic body part to make a movement corresponding to a movement made by the healthy body part when the EMG signals are sensed. Optionally, the controller is configured so that the amplitude of stimulation of at least one of the at least one muscle of the

paretic body part increases when the EMG signal from the corresponding muscle of the healthy body part increases at a corresponding time in the movement of the healthy body part.

In an exemplary embodiment of the invention, the at least one muscle of the paretic body part comprises an antagonistic pair of muscles, and the controller is configured so that the 5 amplitude of stimulation of one muscle of the antagonistic pair of muscles decreases when the EMG signal from the muscle in the healthy body part corresponding to the other muscle of the antagonistic pair of muscles increases at a corresponding time in the movement of the healthy body part.

In an exemplary embodiment of the invention, one or both of the controller and the 10 NMES device are configured so that the stimulation amplitude is not high enough to cause the stimulated muscle to contract in the absence of nerve impulses from the patient's brain, but is high enough to cause the muscle to contract in the presence of nerve impulses from the patient's brain, for at least some patients who cannot move said body part by themselves.

In an exemplary embodiment of the invention, the at least one EMG sensor comprises a 15 plurality of EMG sensors, each EMG sensor adapted to being applied to a different muscle or muscle part of the healthy body part. Optionally, each EMG sensor produces a separate EMG signal. Optionally, the NMES device is adapted to independently stimulate a plurality of muscles or muscle parts of the paretic body part. Optionally, said plurality of muscles or muscle parts of the paretic body part correspond to the muscles or muscle parts of the healthy 20 body part to which the plurality of EMG sensors are adapted to being applied. Optionally, the controller is configured so that amplitude of NMES stimulation of said plurality of muscles or muscle parts of the paretic body part is at least partly dependent on the EMG signals from the plurality of EMG sensors. Optionally, the controller is configured so that the amplitude of NMES stimulation of each of said plurality of muscles or muscle parts depends at least partly 25 on the EMG signal from the corresponding muscle or muscle part.

In an exemplary embodiment of the invention, the paretic body part is a body part that comes in pairs. Optionally, the paretic body part is an arm. Alternatively or additionally, the paretic body part comprises a leg.

In an exemplary embodiment of the invention, the healthy body part belongs to the 30 patient. Alternatively, the healthy body part belongs to a different person.

In an exemplary embodiment of the invention, the controller makes the stimulation amplitude at least partly dependent on a processed form of the EMG signal. Optionally, the processed form of the EMG signal is stretched out in time from the EMG signal. Alternatively or additionally, the processed form of the EMG signal corresponds to an EMG signal that

would be produced by a movement of the healthy body part that is a mirror image of a movement that the healthy part was undergoing when the EMG signal was generated. Alternatively or additionally, the processed form of the EMG signal is time delayed from the EMG signal.

5 In an exemplary embodiment of the invention, the apparatus includes a first position sensing device which monitors a position of the healthy body part. Optionally, the apparatus includes a first actuating device which mechanically changes the position of the healthy body part.

10 In an exemplary embodiment of the invention, the apparatus includes a second position sensing device which monitors a position of the paretic body part.

In an exemplary embodiment of the invention, the apparatus includes a paretic actuating device which mechanically changes the position of the paretic body part under control of said controller and according to said expected move.

15 There is also provided in accordance with an exemplary embodiment of the invention apparatus adapted for rehabilitating a class of patients who have a paretic body part, the apparatus comprising a neuromuscular electrical stimulation (NMES) device adapted to stimulate at least one voluntary muscle in the paretic body, wherein the amplitude of stimulation is not sufficient by itself to cause contraction of said muscle, but the amplitude of stimulation is sufficient to cause contraction of said muscle when a patient in said class attempts to move the body part at the same time.

20 In an exemplary embodiment of the invention, the apparatus comprises at least one paretic EMG sensor adapted for applying to a voluntary muscle of the paretic body part, which at least one paretic EMG sensor produces at least one paretic EMG signal. Optionally, the controller makes the amplitude of stimulation of the paretic body part at least partly dependent 25 on the at least one paretic EMG signal. Optionally, the at least one paretic EMG sensors adapted for applying to the paretic body part comprise a plurality of paretic EMG sensors, each adapted for being applied to a different muscle or muscle part of the paretic body part, and each producing a separate paretic EMG signal. Optionally, the NMES device is adapted to stimulate the muscles or muscle parts of the paretic body that the paretic EMG sensors are adapted for 30 being applied to, and the controller is configured to make the amplitude of stimulation of each muscle or muscle part depend at least partly on the paretic EMG signal from that muscle or muscle part.

In an exemplary embodiment of the invention, said first actuating device mechanically changes the position of said healthy body part at varying levels chosen from the group of complete assistance, partial assistance or no assistance.

5 In an exemplary embodiment of the invention, said first actuating device mechanically changes the position of said healthy body part by limiting the range of motion of said part.

In an exemplary embodiment of the invention, said second actuating device mechanically changes the position of said paretic body part at varying levels chosen from the group of complete assistance, partial assistance or no assistance.

10 In an exemplary embodiment of the invention, said second actuating device mechanically changes the position of said paretic body part by limiting the range of motion of said part.

There is also provided in accordance with an exemplary embodiment of the invention, a method of rehabilitating a patient who has a paretic body part, the method comprising:

- 15 a) having the patient or another person move a healthy body part that is of the same type as the paretic body part;
- b) detecting EMG signals from the healthy body part while it is being moved;
- c) processing said EMG signals to determine at least one property of a NMES signal;
- d) applying a NMES signal to the paretic body part, responsive to said processing; and
- e) moving said paretic body part at most partially by said NMES stimulation.

20 Optionally, said NMES is applied at a timing according to said EMG signals. Alternatively or additionally, said NMES is applied at an amplitude according to said EMG signals.

25 In an exemplary embodiment of the invention, the method comprises having the patient attempt to move the paretic body part, while the NMES is applied, in the same pattern of movement that the healthy body part is moved in while the EMG signals are detected. Optionally, detecting the EMG signals comprises detecting the EMG signals from a plurality of muscles or muscle parts of the healthy body part, and applying NMES comprises applying NMES to a plurality of muscles or muscle parts of the paretic body part corresponding to the plurality of muscles or muscle parts of the healthy body part. Optionally, the amplitude of 30 NMES applied to each muscle or muscle part of the paretic body part during a time interval in the pattern of attempted movement of the paretic body part depends at least partly on the EMG signal detected from the corresponding muscle or muscle part of the healthy body, during a corresponding time interval in the pattern of movement of the healthy body part.

In an exemplary embodiment of the invention, the paretic body part is mechanically provided with movement by an actuating device. Optionally, said actuating device is synchronized to said detected EMG. Alternatively or additionally, said actuating device is synchronized to said applied NMES. Alternatively or additionally, the paretic body part is 5 mechanically assisted with movement by an actuating device. Alternatively or additionally, movement of the paretic body part is limited by an actuating device.

#### BRIEF DESCRIPTION OF THE DRAWINGS

Exemplary embodiments of the invention are described in the following sections with 10 reference to the drawings. The drawings are generally not to scale and the same or similar reference numbers are used for the same or related features on different drawings.

Fig. 1 is a schematic drawing of a healthy arm with EMG electrodes, an EMG unit, a signal processing unit, an NMES unit, and a paretic arm with NMES and EMG electrodes, according to an exemplary embodiment of the invention;

15 Fig. 2 shows plots of a raw EMG signal, rectified signal, smoothed rms signal, according to an exemplary embodiment of the invention;

Fig. 3 is a schematic drawing of an arm attached to a robot arm, according to an exemplary embodiment of the invention;

20 Fig. 4 is a schematic drawing of a paretic arm with NMES electrodes and EMG electrodes, nerve signals from brain, and a signal processing unit, according to a different exemplary embodiment of the invention than Fig. 1; and

Figs. 5A-5G are a time sequence of plots of EMG signals from flexor and extensor signals, at different times during rehabilitation, according to an exemplary embodiment of the invention.

#### DETAILED DESCRIPTION OF EXEMPLARY EMBODIMENTS

25 Fig. 1 shows an apparatus for applying NMES to several muscles of a paretic arm, guided by EMG signals from the corresponding muscles of a healthy arm. Healthy arm 102, belonging either to the patient with the paretic arm or to someone else, has EMG electrodes attached to the skin. The person whose arm it is moves the arm voluntarily in a particular pattern, which generates a certain time-dependent pattern of EMG voltages in the muscles. 30 There are optionally four EMG channels, one channel measuring EMG signals from each of four muscles: the biceps, the triceps, the flexors, and the extensors. Each channel uses three electrodes, two recording signals from near each end of the muscle, and one reference electrode in the middle. For example, electrodes 104 measure the biceps, electrodes 106 measure the triceps, electrodes 108 measure the flexors, and electrodes 110 measure the extensors. It should

be understood from the description of the present invention that similar arrangements are optionally arranged for other muscle pair groups, such as the pectoralis major and deltoid pairs. When the arm is moved voluntarily, these electrodes transmit the EMG signals corresponding to the pattern of muscle contractions producing that movement of the arm, via cable bundles 5 112, 114, 116, and 118, to an EMG device 120. The EMG device, or a separate controller, does preliminary processing of the EMG signals, for example amplifying them, digitizing them, and/or recording them.

The EMG signals are then transmitted, via a cable 122, to a controller 124, which is, for example, a personal computer, or comprises special dedicated hardware. The controller 10 optionally further processes the EMG signals, for example filtering them, rectifying them, smoothing them, changing the timing, or cutting and pasting parts of a sequence in a different order. The signal processing is optionally done automatically, or is partly or entirely under the control of the physical therapist. Fig. 2 shows a plot of a filtered raw EMG signal 202 from one channel, a rectified signal 204, and a smoothed rectified signal 206, in which the root mean 15 square of the signal is calculated in each of a sequence of time intervals, for example every 10 milliseconds. The smoothed rectified signal is a measure of the overall degree of contraction of the muscle or section of the muscle measured by that channel, while eliminating high frequency noise associated with the changes in potential of individual muscle fibers. Optionally, the smoothed rectified EMG signal is averaged over many repetitions of the same 20 pattern of movement. Optionally, the EMG sensors add up or average the EMG signals coming from several different sections of the muscle, or this is done by the signal processing. An optional type of signal processing is validating EMG signals, for example, by measuring the actual motion of a healthy limb while measuring EMG thereon, so that it is clear that the EMG signals result from the desired movement. Optionally, one or more of the parameters of the 25 signal processing are controlled by the therapist. In some embodiments of the invention, the smoothed rectified EMG signal is created by dispensing with anomalous EMG readings, such as not factoring the high and low readings, and averaging the rest of the readings over multiple repetitions. In some embodiments, the EMG is measured and the NMES applied, at a time delay, for example, to allow multiple EMG signals to be collected and averaged.

30 Controller 124 also controls NMES device 126, via cable 128. Optionally, controller 124 commands NMES device to produce NMES signals in each of four channels. The signals in the four channels travel through cables 130, 132, 134, and 136, to electrodes 138, 140, 142, and 144, which respectively stimulate the biceps, triceps, flexors, and extensors on the patient's paretic arm 146 (or optionally any other muscle pair as the case may be). Optionally, the

NMES signals in each channel are given a time-dependent amplitude which will produce the same movement in the paretic arm as was performed by the healthy arm. This is done, for example, by making the signal strength in each NMES channel depend on the processed signal amplitude from a corresponding one of the four EMG channels. For example, the NMES signal 5 is proportional to the processed EMG signal amplitude, or is a fixed monotonic function of the processed EMG signal amplitude, for the corresponding channel.

Optionally, the NMES signal depends also on the EMG signal from one or more other channels. For example, because the biceps and triceps work against each other, the NMES signals controlling the biceps and triceps, optionally depend on a linear combination of the 10 EMG signal from the biceps and the EMG signal from the triceps, with a negative coefficient for the EMG signal from the triceps. If the linear combination is positive, only the biceps is stimulated, and if the linear combination is negative, only the triceps is stimulated. A similar method is optionally used for any other pair of agonist antagonist muscles, such as the flexors and extensors which also work against each other.

15 Optionally, the NMES signals are based not directly on the EMG signals from the corresponding muscles, but are modified to produce motion that is reversed in some way from the motion associated with the EMG signals. For example, if the EMG signals come from a left arm and the NMES signals are applied to a right arm, then optionally the NMES signals are changed to produce motion in the right arm that is the same as the motion of the left arm, rather 20 than a mirror image of it, as would occur if the corresponding muscles in the two arms were to contract at the same time. Alternatively or additionally, whether or not the two arms are a left arm and a right arm, if the motion of the healthy arm is cyclical, then the NMES signals are changed to produce motion in the paretic arm that is 180 degrees out of phase from the motion 25 of the healthy arm. Such a modification in the NMES signals might be particularly useful to use for the left and right legs, for example, in a patient who needs to relearn how to walk or ride a bike.

30 Optionally, there are also EMG electrodes 148, 150, 152, and 154, attached to the paretic arm. These sensors send signals along cables 156, 158, 160 and 162, respectively, to four additional channels of EMG device 120, which thus has a total of eight channels. These additional EMG signals are processed by the EMG device and by controller 124, similar to the processing of the EMG signals from healthy arm 102. Optionally, the EMG signals from paretic arm 146 are also used by controller 124 in controlling the NMES signals. The EMG signals in paretic arm 146 may arise because the sensory-motor cortex of the patient is still capable of producing weak nerve impulses in paretic arm 146, even if these nerve impulses are

too weak to cause the paretic arm to move. By timing the NMES signals to the corresponding EMG signals in the paretic arm, the paretic arm can move in response to the attempts of the patient to move it, providing kinesthetic feedback to the patient. Alternatively or additionally, EMG signals in paretic arm 146 may be induced by moving paretic arm 146 passively, for 5 example by a robot arm as discussed below in describing Fig. 3.

Optionally, controller 124 also uses other information in controlling the strength of the NMES signals. For example, the healthy arm has a sensor 164, for example a strain sensor, which measures the degree of bending of the elbow, and a sensor 166 which measures the degree of extension of the fingers, while the paretic arm has similar sensors 168 and 170. The 10 sensors feed into a unit 172 which processes the sensor data to determine the bending of the arm and fingers, and this information is conveyed, for example by cable 174, to controller 124. Optionally, unit 172 and controller 124 are part of a single control unit. Optionally, sensors are used only with one of the arms. Optionally there are other sensors which measure other aspects of the arm and hand position, particularly if EMG and NMES is used with additional muscles. 15 A variety of other types of sensors are additionally or alternatively used for measuring the arm or hand position, for example the arm is fitted to a robot arm which has such sensors to measure its own state, as shown in Fig. 3 which will be described below. Or, LEDs are attached to key points on the arm and hand and their location tracked with a video camera, or magnetic field sensors are attached to key points on the arm and hand, and an external 20 magnetic field and/or field gradient imposed. Other methods will be apparent to those skilled in the art of sensing the position and orientation of body parts.

The position of the paretic arm can be used, for example, as negative feedback to the NMES signals. During the course of rehabilitation, as the patient's own nerve impulses become stronger and/or more effective, for example distinguishing better between antagonistic pairs of 25 muscles, the NMES signal can be reduced while producing the same arm motion. This kind of feedback can also be used within a given rehabilitation session. For example, if the patient is momentarily having trouble continuing to move his arm, the NMES amplitude is momentarily increased, until the patient is able to start moving his arm again. Optionally, in this case, the controller distinguishes between the patient simply resting, and the patient trying 30 unsuccessfully to move his arm, for example by looking at EMG signal levels in the paretic arm.

The position of the healthy arm can be used, for example, to supplement the EMG signals from the healthy arm, as a measure of the degree of contraction of the muscles in the

healthy arm. Alternatively or additionally, the data on position of both arms can be used to monitor the progress of the rehabilitation of the patient.

Fig. 3 shows an arm 302, which could be either the healthy arm or the paretic arm in Fig. 1, attached to a robot arm 300. The upper arm is held by a holder 304, and the lower arm is held by a holder 306. Upper arm holder 304 is attached to an extendable rod 308, which is connected to a controllable ball joint 310, and similarly lower arm holder 306 is attached to an extendable rod 312, which is connected to a controllable ball joint 314. Ball joints 310 and 314 are connected to each other with a rigid connector 316. The ball joints and extendable rods include both actuators and sensors, for all their degrees of freedom, in this case two degrees of freedom for each ball joint, and one degree of freedom for each extendable rod. The sensors can sense the degree of bending of the elbow of arm 302, and the actuators can apply force to bend or unbend the elbow, and/or to resist bending or unbending of the elbow by the patient. Optionally, the actuators and sensors have more or fewer degrees of freedom, depending for example on which muscles are being rehabilitated, and optionally the robot arm is attached to additional points on arm 302, for example to different points on the wrist, hand, and fingers. Signals from the sensors and to the actuators are processed by a robot arm control box, not shown in Fig. 3.

The robot arm optionally is used in the same way as sensors 164, 166, 168 and 170 in Fig. 1. In addition, the fact that the robot arm can move in a controlled way under its own power means that it can supplement the NMES, in providing kinesthetic feedback for the patient, if used with the paretic arm. The robot arm provides a different kind of kinesthetic feedback than NMES provides, since it moves the paretic arm without causing muscles to contract under their own power, and both types of feedback are potentially useful in rehabilitation. For example, NMES may not be able to produce a smooth and accurate motion by itself, and robot arm can help to correct and smooth the motion induced by NMES.

Optionally, both the passive and active modes of the robot arm are combined with the NMES. Movement generated by the robot arm is assisted by contraction of the muscles by NMES. When the patient moves the robot arm in an active way, the NMES signals are adjusted correspondingly.

30 Optionally, motion of the robot arm attached to the paretic arm is based on EMG readings in combination with motion of the healthy arm.

When providing rehabilitation various types of motion may be supported by the robotic arm, for example, one or more of:

- a) Passive motion. The robot arm is moved and the patient moves with it.

b) Resisted motion. The patient moves the robot arm and encounters resistance. The resistance may be of various magnitudes and may be uniform in all direction or be directional.

c) Assisted motion. When a patient moves the robot arm, a positive feedback on arm increases the force of motion in the direction moved by the patient.

5 d) Force field motion. The patient moves the robot arm. Along a certain trajectory one level of resistance (or none) is encountered. Deviation from the trajectory is not allowed or meets with resistance. Motion along a "correct" trajectory can be without resistance, or possibly assisted. An increased resistance is exhibited in a volume surrounding trajectory. An even greater resistance is exhibited in a surrounding volume. A prevention of motion may be provided in an outside volume. In an exemplary embodiment of the invention, a corrective force vector is applied when not on trajectory, pointing towards trajectory. Optionally, instead of a corrective force, resistance varies as a function of distance from trajectory, thus, motion of the robot arm is naturally urged back to trajectory. Optionally, the force is applied in the direction of the path. Alternatively, the force maybe a unidirectional force of resistance.

10 15 This type of motion helps to train the patient in a desired motion.

e) Mirrored motion. Motion of the robot arm is required to mirror the trajectory of motion of a different element, for example for dual limb rehabilitation.

20 f) Free motion. Patient moves the robot arm in any way he desires, possibly receiving feedback. As the patient (or therapist or helper) moves the robot arm, a device may record it for future playback. In a playback mode the prerecorded motion (or path) is optionally reconstructed using other modes. Optionally, the recorded path is modified (e.g., smoothed), for example automatically or manually.

25 g) General Force Field. A force field and/or an assistance field is defined which is not related to any particular trajectory. For example, a range of trajectories may be allowed to be practiced by a user, or a real or virtual situation simulated (e.g., water, areas with obstacles).

h) Local force field. A force field which is applied to only a small locality and/or only in one or two dimensions.

30 i) Restricted motion. One or more points of the body of a subject are supported or prevented from moving. Optionally, the angles between such points and the moving points on the device are measured. In one example the elbow is locked with a dedicated harness allowing only a shoulder motion. In some embodiments, the restriction is partial and/or is provided by a movable element (e.g., an arm).

j) Initiated Motion. The patient initiates the motion (e.g., a 1 cm motion or 100 gram force) and the robot arm completes or helps the patient complete the motion in space. The completion may be of a whole trajectory or of part of a trajectory.

5 k) Implied motion. The robot arm begins the motion and the patient completes it. The robot arm may assist the rest of the motion in various manners (e.g., by changing to one of the modes described herein after the motion starts). If the patient fails to pick up the motion, the robot arm may generate a cue, for example an audio reminder. Different parts of a single motion trajectory may each have a machine initiation definition. Optionally, if a patient is too slow in moving, the robot arm begins the motion.

10 l) Cued motion. The patient receives a cue from the system before motion according to a different mode begins. The cue can be, for example, vibration of the robot arm, stimulation pads on the skin, audio or visual cue. In some embodiments of the invention, the strength of the cue and/or its timing and/or other ongoing activities (e.g., a visual display and game) are used to help train the coordination between different modalities, for example, hand-eye 15 coordination. A motion cue can be used to train a kinesthetic sense.

20 m) Teach Mode. The robot arm is taught a motion. In one example, a therapist performs a motion and motion parameters at each point are recorded and can then be used for an exercise. Another way of teaching the system is to follow a path that the therapist traces with his/her own motion. The therapist may use the controller to indicate a point to be taught or a continuous mode may be defined by which an entire trajectory is learned. Optionally the path and points are edited before replay. Optionally, the paths are processed, for example, by smoothing or identifying motion points, before playback.

Thus, in some embodiments of the invention, the robot arm can provide one or more of Isokinetic, Isotonic and Isostatic exercises.

25 It should be appreciated that a definition of a trajectory which the robot arm is to follow can include speed parameters. For example, a user may be assisted, or urged, or expected, to move the robot arm at a certain speed. The speed may be, for example, absolute, or relative (e.g., requiring a uniform speed or the speed to match a non-uniform profile).

30 Optionally, an angular trajectory is defined, which places constraints on an angular orientation of the robot arm. In some embodiments, the constraint is one dimensional. In others it is two or three dimensional.

Speed, angles and spatial trajectories in a particular rehabilitation scenario may each belong to a different one of the above motion types. For example, spatial trajectory may be of a force field type, while speed trajectory is free or assisted. The type of trajectory and/or its

parameters may also vary along the trajectory, as a function of time and/or as a function of previous performance. For example, a smaller assistance at a later part of a trajectory may be provided for a type of motion which was properly (or better than expected) executed in an earlier part of the trajectory.

5        Trajectories may be absolute, for example, defined as a function of a resting point or a different point on the robot arm. In other embodiments, the trajectories are purely relative, for example, requiring a patient to move an arm in a straight line, regardless of starting point. In other embodiments, a trajectory is partially relative, in that once motion starts, this determines the shape of the rest of the trajectory, for example, a start of a trajectory indicating if a patient  
10      is standing or sitting, and thus what type of hand motion is expected.

In some embodiments, such as described below, where multiple points are defined, the trajectories of each point may be of different types. In some embodiments, what is defined is a trajectory as a function of two or more points. For example, if two points are used to define an elbow configuration (e.g., angle between bones), the trajectory constraints may be defined on  
15      the motion of the elbow. Such motion may be relative in space (e.g., a comparison of the two points) and not absolute (e.g., compared to a device reference point). It should be noted that in some embodiments of the invention a tensor or tensor field is provided, as each point in space can have associated with it a speed, a force and/or a rotation, all of which can be scalar or a vector.

20      In some embodiments of the invention, different modes are defined for different parts of a trajectory or for different parts of space (e.g., for a particular arm). Optionally, a mode may be triggered based on the actual performance. For example, if motion velocity is below a certain threshold, a more assistive mode is provided. Similarly, a pause of over a threshold may imply a more assistive mode. An exact motion may imply a less assistive mode.

25      Fig. 4 shows an arrangement according to another exemplary embodiment of the invention, in which only a paretic arm 146 is used. As in Fig. 1, there are EMG electrodes 148, 150, 152 and 154, attached respectively to the biceps, triceps, flexors and extensors of the paretic arm, three electrodes for each muscle, and the EMG signals are conveyed along cables 156, 158, 160, and 162, to an EMG device 120, which, after preliminary processing, transmits  
30      the signals to controller 124. As in Fig. 1, controller 124 uses the EMG signals in determining the amplitude and timing of NMES signals transmitted by NMES device 126, which stimulate the biceps, triceps, flexors and extensors of paretic arm 146 through NMES electrodes 138, 140, 142, and 144.

The NMES signals transmitted by NMES device 126 in Fig. 4 are not strong enough, in themselves, to cause paretic arm 146 to move significantly, and this is optionally also true in Fig. 1. But the NMES signals, together with the patient's own nerve impulses, are strong enough to cause the arm to move. Thus, the paretic arm only moves when the patient tries to move it, and the kinesthetic feedback provided by the motion further encourages the development of alternate pathways for nerve impulses in the patient, or alternate locations in the motor cortex to originate nerve impulses to the same muscles, eventually enabling the patient to move the paretic arm by himself. This may be particularly useful when the motion involves a coordinated sequence of contractions of more than one muscle. Optionally, in the course of rehabilitation, the NMES signal is lowered, as less NMES signal is needed in order to allow the patient to move the paretic arm.

Optionally, in the absence of nerve impulses from the patient's motor cortex, the NMES signals are between 100% and 120% of the amplitude needed to produce motion for an average healthy subject, or for an average paretic patient, or they are adjusted to that level for a particular patient. Alternatively, they are between 120% and 140% of that amplitude for any of these people, or between 80% and 100%, or between 60% and 80%, or less than 60%. Optionally, for any of these people, the NMES signals are between 100% and 120% of the level needed to produce motion in the presence of nerve impulses from the motor cortex when the person makes an effort to move, or between 120% and 140%, or between 140% and 200%, or greater than 200%.

Optionally, the NMES is targeted to a part of the muscle with at least as much spatial precision as an average healthy subject is able to achieve when voluntarily directing nerve impulses to that muscle. Alternatively, the NMES is targeted with less than this much precision, but with at least half this much precision, or with less than half this much precision, but at least one quarter this much precision, or with less than one quarter this much precision.

Characteristics of the procedure illustrated in Figs. 1 and 4 can be varied to adapt to the needs of the patient, in order to facilitate rehabilitation. Several examples are given below.

The EMG and NMES need not use the four muscles shown being used in Figs. 1 and 4, but could include more muscles, or fewer muscles. Optionally, only the biceps and triceps are used initially. Then, once the patient has gained some ability to use the biceps and triceps effectively, the flexors and extensors are added to the EMG and NMES channels. These four muscles are basic to gross control of the arm, in addition to other muscles such as the pectorals and the deltoids. Later, individual fingers are added, and/or other wrist and hand motions, to

improve fine motor control. For rehabilitation of body parts other than the arm, of course, other groups of muscles are selected.

The amplitude of NMES optionally varies depending on feedback from various sources, and depending on the immediate goal of the rehabilitation program. As mentioned previously, 5 the NMES signal is optionally decreased as the patient recovers the ability to generate nerve impulses and move his muscles by himself. Alternatively, if the immediate goal is the strengthening of atrophied muscles, the amplitude of NMES is optionally increased as the muscle gets stronger, and can benefit from more strenuous exercise. In this case, the arm is 10 optionally made to move against a restraining force, for example a weight or a robotic arm, which is increased as the muscle gets stronger, so a stronger NMES signal is required in order to move the arm by the same amount.

In addition to using kinesthetic feedback to encourage the development of alternative pathways for nerve impulses, as discussed above, other kinds of feedback are optionally used to help the patient learn how to control his muscles more effectively. For example, seeing the 15 movement of the arm, when his nerve impulses are supplemented by NMES stimulation, can help the patient adjust his efforts to move his arm. Similarly, such feedback for conscious learning by the patient can be provided by a device, such as the robot arm in Fig. 3, which measures and records the motion of the arm, and by the processed EMG signals. For example, the patient can try to make the EMG signals from the paretic arm more closely resemble the 20 EMG signals generated by the healthy arm when it is performing the desired movement, or he can try to make the EMG signals from the paretic arm more closely resemble some template, perhaps developed from examining recorded EMG signals from the healthy arm, or from paretic arms of other patients who have undergone similar rehabilitation.

In the arrangement shown in Fig. 1, if the patient's other arm is used as the healthy arm, 25 then optionally the patient tries to move both arms synchronously, in mirror image movements. The NMES signals, optionally based on the EMG signals of the healthy arm, allow the paretic arm to move, and since the patient is attempting to move both arms in synchrony, he receives kinesthetic feedback from the paretic arm, which helps promote the development of alternate pathways for nerve impulses. In some embodiments of the present invention, movement of the 30 paretic arm to mirror movement of the healthy arm is assisted, in part or in whole, by the robotic arm 300.

Optionally, the NMES signals are adapted to the capability of the paretic arm. For example, if the muscles in the paretic arm are incapable of responding as rapidly as normal to the NMES, then the NMES signals are optionally slowed down, or high frequency components

are reduced or removed. As the muscles recover the capability of more rapid response, the NMES signals are sped up again. The speed of the NMES signals is either adjusted automatically, in response to sensor data on movement of the paretic arm, or manually by the therapist, optionally using such sensor data to evaluate the patient. If a robotic arm is used in coordination with NMES to help move the paretic arm, the motion of the robotic arm is optionally slowed down together with the NMES. Even if the robotic arm is used to help move the paretic arm without NMES, the motion of the robotic arm is optionally slowed down if, for example, this will help the patient to make a greater contribution to the motion with his own nerve impulses, or will be useful for some other reason in rehabilitation.

10 Figs. 5A through 5G illustrate a procedure for rehabilitating a patient who has a problem that is common following a stroke in the somato-motor cortex—the failure of the patient's nerve impulses to distinguish adequately between two muscles that form an antagonistic pair, such as the biceps and triceps, or the flexors and extensors. As shown in Fig. 5A, the EMG signal 402 from the flexors, and signal 404 from the extensors, when the patient attempts to open and close her hand, are strong enough to cause both muscles to contract, since they are above threshold level 406. But both muscles contract at the same time, so that they work against each other, and the hand exhibits very little movement. First, the patient learns to decrease the overall activity of both the flexors and extensors, below the threshold for contraction, as shown progressively in Figs. 5B, 5C, and 5D. Then, as shown in Figs. 5E, 5F, and 5G, the patient is taught to increase the activity of the extensors, while keeping the flexors relaxed. This is done, for example, by applying NMES to the extensors, increasing kinesthetic feedback, when patient tries to contract the extensors.

To summarize, listed below are some of the rehabilitation methods that can be used in some embodiments of the invention, including those discussed.

25 1) Record EMG in healthy arm and apply similar pattern of NMES to paretic arm, in real time or not.

2) Adjust NMES amplitude to supplement nerve impulses in paretic arm, as measured by EMG in paretic arm.

3) Target NMES to sections of paretic arm where EMG is weak.

30 4) Slow down NMES to adapt to slow response time of paretic arm.

5) Have patient move both arms together, in mirror image, while applying NMES based on EMG in healthy arm.

- 6) Have patient move both arms together not in a mirror image, and/or in a cyclical motion 180 degrees out of phase, while applying NMES based on (but modified from) EMG in healthy arm.
- 7) Base NMES on average EMG over many repetitions of movement by healthy arm.
- 5 8) Sense position of paretic arm and use negative feedback for NMES; optionally use EMG of paretic arm to distinguish inability to move arm from intentional resting.
- 9) Record sensed position of healthy arm as a function of time while recording EMG signals, then apply corresponding NMES to paretic arm when paretic arm is in a corresponding position.
- 10 10) Use robotic arm to move healthy arm in a desired pattern, detect the resulting EMG signals generated passively in the healthy arm, and use them as a basis for NMES applied to paretic arm to produce corresponding motion.
- 11) Use robotic arm and/or NMES to move or assist moving paretic arm, matching to measured position of healthy arm.
- 15 12) Use robot arm to measure resistance of paretic arm to motion, thereby determining whether failure of paretic arm to move is due to failure of muscle to contract, or failure to differentiate between antagonistic pairs of muscles; optionally adjust NMES accordingly.
- 13) Use robotic arm, with or without NMES, to assist moving paretic arm, slowing down robotic arm to match capability of paretic arm.
- 20 14) Use robotic arm to work against muscles of paretic arm, with or without NMES, optionally adapting force to capability of paretic arm.
- 15) Use EMG of paretic arm to teach patient to better control paretic arm, optionally including better differentiating between antagonistic pairs of muscles.

In an exemplary embodiment of the invention, the method of applying NMES is combined with the teachings of other applications filed by the applicant.

US provisional application no. 60/633,442 filed on December 7, 2004, also being filed as PCT application on same date as the present application and by the same applicant, entitled "Methods and Apparatuses for Rehabilitation Exercise and Training" and having attorney docket number 414/04388, the disclosures of which are incorporated herein by reference, 30 describe rehabilitation of balance. In an exemplary embodiment of the invention, NMES is used to help a patient coordinate two sides of a body, or to stimulate muscle sin one part of the body relating to balance (e.g., a torso) when another part moves (e.g., legs).

US provisional application no. 60/566,079 filed on April 29, 2004, also being filed as PCT application on same date as the present application and by the same applicant, entitled

“Fine Motor Control Rehabilitation” and having attorney docket number 414/04401, the disclosures of which are incorporated herein by reference, describe rehabilitation of fine motor control. In an exemplary embodiment of the invention, NMES is used to help a patient coordinate gross and fine motor motions (e.g., measuring a large muscle and stimulating a small muscle or vice versa) or to copy fine motor control from a healthy arm to a paretic arm.

5 US provisional application no. 60/633,428 filed on December 7, 2004, also being filed as PCT application on same date as the present application and by the same applicant, entitled “Gait Rehabilitation Methods and Apparatuses” and having attorney docket number 414/04391, the disclosures of which are incorporated herein by reference, describe 10 rehabilitation of gait. In an exemplary embodiment of the invention, NMES is used to coordinate the movement of two legs and/or of motion of different parts of a leg, for example, EMG measurements on a thigh (healthy or paretic) are used to drive NMES signals to a paretic calf.

15 US provisional application no. 60/542,022 filed on February 5, 2004, also being filed as PCT applications on same date as the present application and by the same applicant, entitled “Methods and Apparatus for Rehabilitation and Training”, the disclosures of which are incorporated herein by reference, describe rehabilitation devices of various types. In an exemplary embodiment of the invention, NMES stimulation and/or EMG measurement is provided in such devices.

20 US provisional application no. 60/604,615 filed on August 25, 2004, the disclosure of which is incorporated herein by reference, describes rehabilitation while measuring and/or otherwise taking into account brain plasticity. In an exemplary embodiment of the invention, NMES stimulation is provided in a manner which is coordinated with the activity of brain 25 centers responsible for generating neural signal to the paretic limb and/or to receive signals from the paretic limb. Such a brain area may be detected, for example using EEG or fMRI methods.

30 As used herein, a “position” of an arm or another body part may include not just the spatial location of a particular portion of the arm or body part, but any other information needed to specify its spatial state, including, for example, how much it is bent at the elbow, how much the forearm is twisted, how much the wrist is bent, etc. In some embodiments, the velocity of the part and/or its orientation are controlled.

The invention has been described in the context of the best mode for carrying it out. It should be understood that not all features shown in the drawing or described in the associated text may be present in an actual device, in accordance with some embodiments of the

invention. Furthermore, variations on the method and apparatus shown are included within the scope of the invention, which is limited only by the claims. Also, features of one embodiment may be provided in conjunction with features of a different embodiment of the invention. As used herein, the terms "have", "include" and "comprise" or their conjugates mean "including but not limited to."

CLAIMS

1. Apparatus for rehabilitating a patient who has a paretic body part, the apparatus comprising:

5 a) at least one electromyography (EMG) sensor adapted to being applied to a voluntary muscle of a healthy body part of the same type as the paretic body part, which at least one sensor produces at least one EMG signal;

b) a neuromuscular electrical stimulation (NMES) device adapted for stimulating at least one voluntary muscle of the paretic body part;

10 c) a controller which controls the NMES device and expects a motion of the paretic body part, making the amplitude of stimulation of the paretic body part at least partly dependent on the EMG signal from the healthy body part such that said NMES stimulation is not sufficient, on its own, to move said paretic body part said expected motion.

15

2. Apparatus according to claim 1, wherein the at least one muscle of the healthy body part corresponds to the at least one muscle of the paretic body part.

3. Apparatus according to claim 1, wherein said controller is configured to process said  
20 EMG signals and determine at least one property of said NMES signal.

4. Apparatus according to claim 2, wherein the controller is configured so that the NMES stimulates the paretic body part to make a movement corresponding to a movement made by the healthy body part when the EMG signals are sensed.

25

5. Apparatus according to claim 4, wherein the controller is configured so that the amplitude of stimulation of at least one of the at least one muscle of the paretic body part increases when the EMG signal from the corresponding muscle of the healthy body part increases at a corresponding time in the movement of the healthy body part.

30

6. Apparatus according to claim 4 or claim 5, wherein the at least one muscle of the paretic body part comprises an antagonistic pair of muscles, and the controller is configured so that the amplitude of stimulation of one muscle of the antagonistic pair of muscles decreases when the EMG signal from the muscle in the healthy body part corresponding to the other

muscle of the antagonistic pair of muscles increases at a corresponding time in the movement of the healthy body part.

7. Apparatus according to any of the preceding claims, wherein one or both of the controller and the NMES device are configured so that the stimulation amplitude is not high enough to cause the stimulated muscle to contract in the absence of nerve impulses from the patient's brain, but is high enough to cause the muscle to contract in the presence of nerve impulses from the patient's brain, for at least some patients who cannot move said body part by themselves.

10

8. Apparatus according to any of the preceding claims, wherein the at least one EMG sensor comprises a plurality of EMG sensors, each EMG sensor adapted to being applied to a different muscle or muscle part of the healthy body part.

15

9. Apparatus according to claim 8, wherein each EMG sensor produces a separate EMG signal.

10. Apparatus according to claim 9, wherein the NMES device is adapted to independently stimulate a plurality of muscles or muscle parts of the paretic body part.

20

11. Apparatus according to claim 10, wherein said plurality of muscles or muscle parts of the paretic body part correspond to the muscles or muscle parts of the healthy body part to which the plurality of EMG sensors are adapted to being applied.

25

12. Apparatus according to claim 11, wherein the controller is configured so that amplitude of NMES stimulation of said plurality of muscles or muscle parts of the paretic body part is at least partly dependent on the EMG signals from the plurality of EMG sensors.

30

13. Apparatus according to claim 12, wherein the controller is configured so that the amplitude of NMES stimulation of each of said plurality of muscles or muscle parts depends at least partly on the EMG signal from the corresponding muscle or muscle part.

14. Apparatus according to any of the preceding claims, wherein the paretic body part is a body part that comes in pairs.

15. Apparatus according to claim 14, wherein the paretic body part is an arm.

16. Apparatus according to claim 14, wherein the paretic body part is a leg.

5

17. Apparatus according to any of claims 14-16, wherein the healthy body part belongs to the patient.

18. Apparatus according to any of claims 14-16, wherein the healthy body part belongs to a  
10 different person.

19. Apparatus according to any of the preceding claims, wherein the controller makes the stimulation amplitude at least partly dependent on a processed form of the EMG signal.

15 20. Apparatus according to claim 19, wherein the processed form of the EMG signal is stretched out in time from the EMG signal.

21. Apparatus according to claim 19 or claim 20, wherein the processed form of the EMG signal corresponds to an EMG signal that would be produced by a movement of the healthy  
20 body part that is a mirror image of a movement that the healthy part was undergoing when the EMG signal was generated.

22. Apparatus according to any of claims 19-21, wherein the processed form of the EMG signal is time delayed from the EMG signal.

25

23. Apparatus according to any of the preceding claims, also including a first position sensing device which monitors a position of the healthy body part.

30 24. Apparatus according to claim 23, also including a first actuating device which mechanically changes the position of the healthy body part.

25. Apparatus according to any of the preceding claims, also including a second position sensing device which monitors a position of the paretic body part.

26. Apparatus according to any of claims 1-25, including a paretic actuating device which mechanically changes the position of the paretic body part under control of said controller and according to said expected move.

5 27. Apparatus according to claim 24, wherein said first actuating device mechanically changes the position of said healthy body part at varying levels chosen from the group of complete assistance, partial assistance or no assistance.

10 28. Apparatus according to claim 24, wherein said first actuating device mechanically changes the position of said healthy body part by limiting the range of motion of said part.

29. Apparatus according to claim 26, wherein said second actuating device mechanically changes the position of said paretic body part at varying levels chosen from the group of complete assistance, partial assistance or no assistance.

15 30. Apparatus according to claim 26, wherein said second actuating device mechanically changes the position of said paretic body part by limiting the range of motion of said part.

31. Apparatus adapted for rehabilitating a class of patients who have a paretic body part, 20 the apparatus comprising a neuromuscular electrical stimulation (NMES) device adapted to stimulate at least one voluntary muscle in the paretic body, wherein the amplitude of stimulation is not sufficient by itself to cause contraction of said muscle, but the amplitude of stimulation is sufficient to cause contraction of said muscle when a patient in said class attempts to move the body part at the same time.

25 32. Apparatus according to any of the preceding claims, also including at least one paretic EMG sensor adapted for applying to a voluntary muscle of the paretic body part, which at least one paretic EMG sensor produces at least one paretic EMG signal.

30 33. Apparatus according to claim 32, wherein the controller makes the amplitude of stimulation of the paretic body part at least partly dependent on the at least one paretic EMG signal.

34. Apparatus according to claim 33, wherein the at least one paretic EMG sensors adapted for applying to the paretic body part comprise a plurality of paretic EMG sensors, each adapted for being applied to a different muscle or muscle part of the paretic body part, and each producing a separate paretic EMG signal.

5

35. Apparatus according to claim 34, wherein the NMES device is adapted to stimulate the muscles or muscle parts of the paretic body that the paretic EMG sensors are adapted for being applied to, and the controller is configured to make the amplitude of stimulation of each muscle or muscle part depend at least partly on the paretic EMG signal from that muscle or muscle part.

10

36. A method of rehabilitating a patient who has a paretic body part, the method comprising:

- a) having the patient or another person move a healthy body part that is of the same type as the paretic body part;
- b) detecting EMG signals from the healthy body part while it is being moved;
- c) processing said EMG signals to determine at least one property of a NMES signal;
- d) applying a NMES signal to the paretic body part, responsive to said processing; and
- e) moving said paretic body part at most partially by said NEMS stimulation.

20

37. A method according to claim 36, wherein said NMES is applied at a timing according to said EMG signals.

25

38. A method according to claim 36, wherein said NMES is applied at an amplitude according to said EMG signals.

39. A method according to claim 36, also including having the patient attempt to move the paretic body part, while the NMES is applied, in the same pattern of movement that the healthy body part is moved in while the EMG signals are detected.

30

40. A method according to claim 39, wherein detecting the EMG signals comprises detecting the EMG signals from a plurality of muscles or muscle parts of the healthy body part, and applying NMES comprises applying NMES to a plurality of muscles or muscle parts of the

paretic body part corresponding to the plurality of muscles or muscle parts of the healthy body part.

41. A method according to claim 40, wherein the amplitude of NMES applied to each muscle or muscle part of the paretic body part during a time interval in the pattern of attempted movement of the paretic body part depends at least partly on the EMG signal detected from the corresponding muscle or muscle part of the healthy body, during a corresponding time interval in the pattern of movement of the healthy body part.

10 42. A method according to any of claims 36-41, wherein the paretic body part is mechanically provided with movement by an actuating device.

43. A method according to claim 42, wherein said actuating device is synchronized to said detected EMG.

15 44. A method according to claim 42, wherein said actuating device is synchronized to said applied NMES.

45. A method according to any of claims 36-44, wherein the paretic body part is 20 mechanically assisted with movement by an actuating device.

46. A method according to any of claims 36-45, wherein movement of the paretic body part is limited by an actuating device.

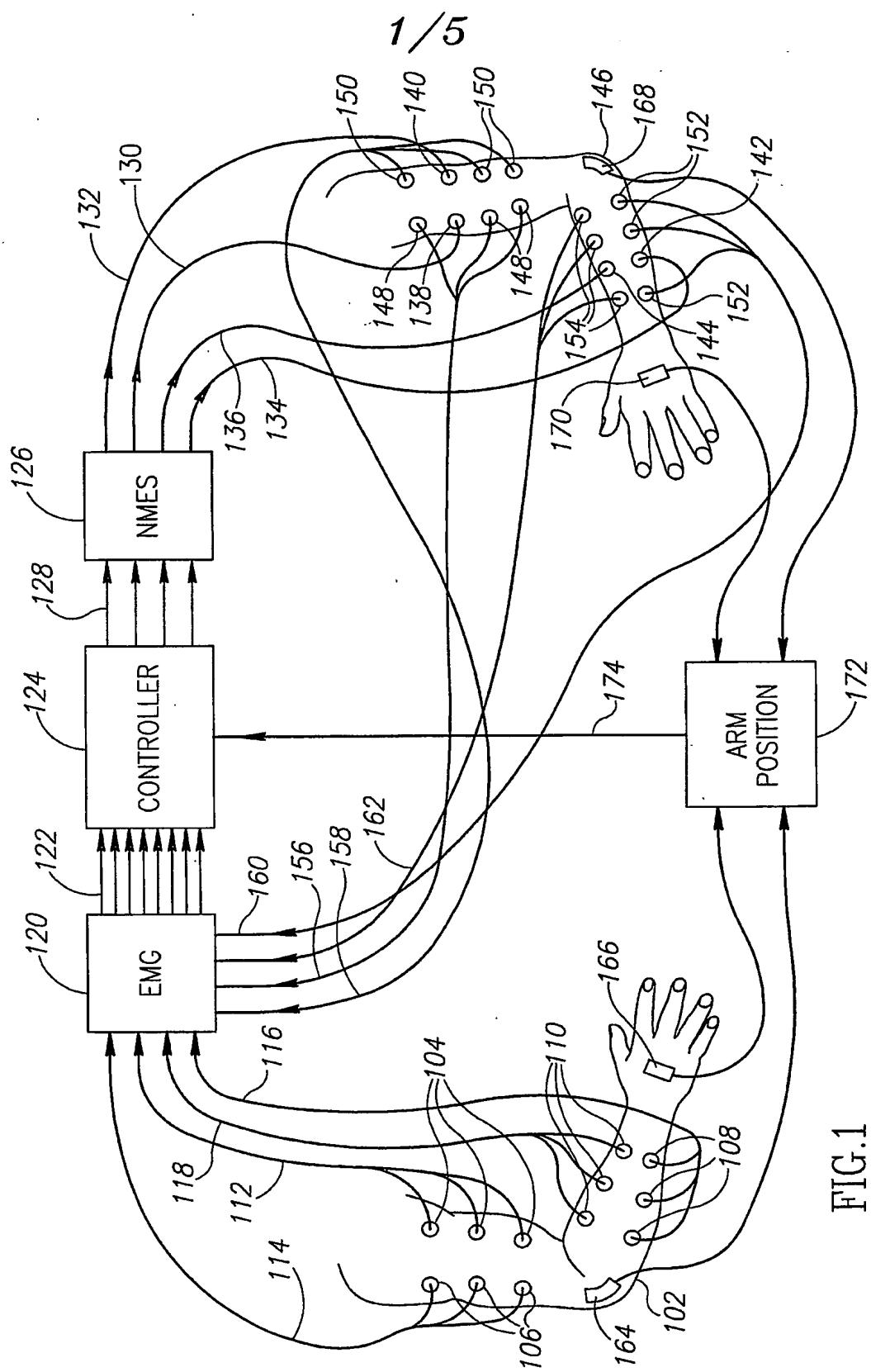


FIG.1

2/5

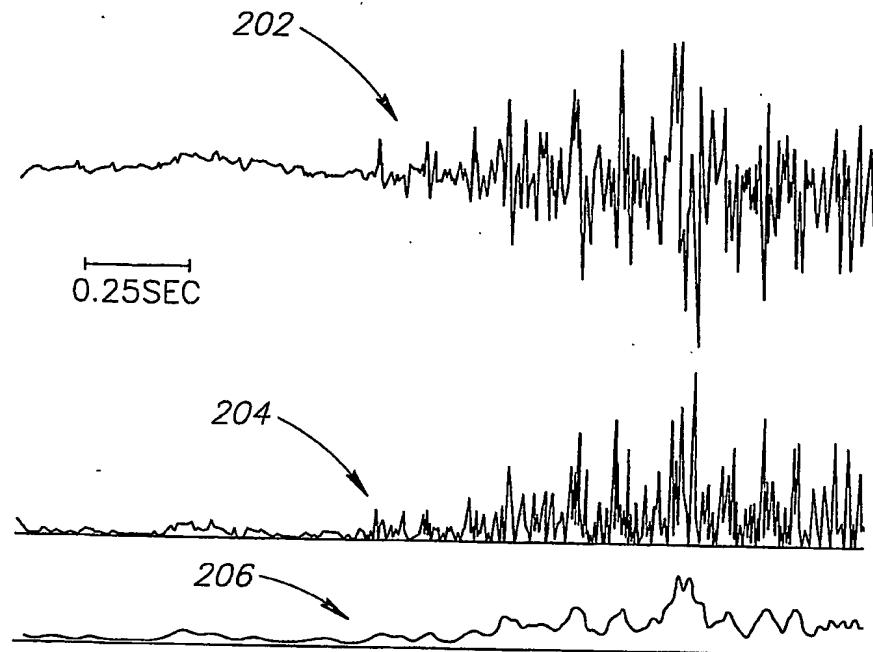


FIG.2

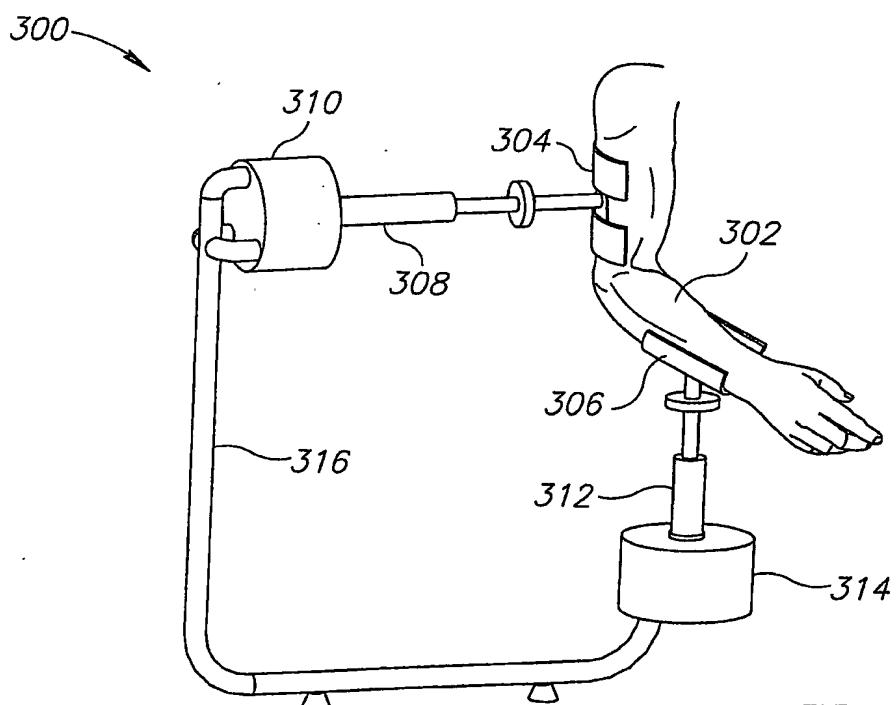


FIG.3

3/5

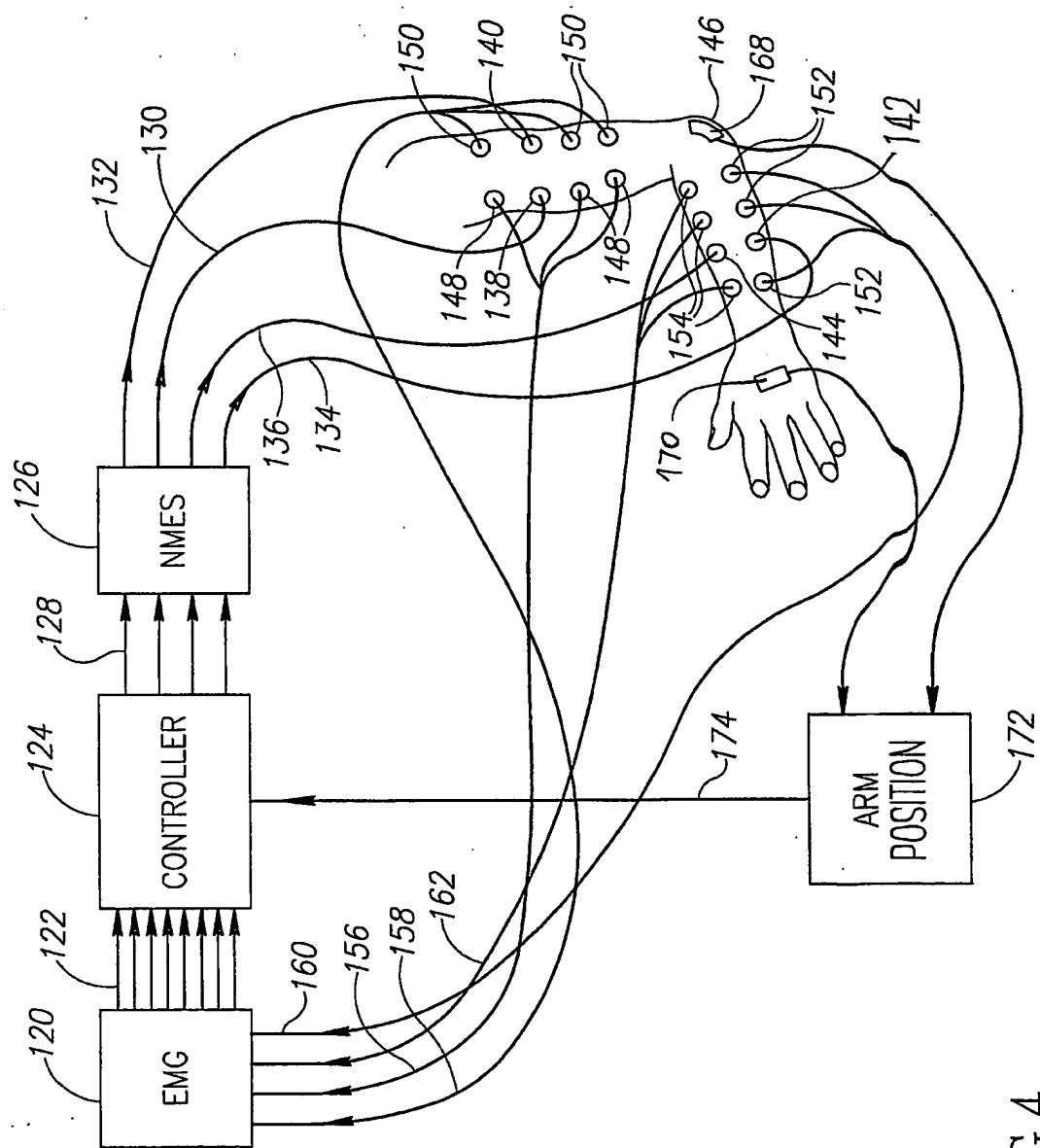


FIG.4

4/5

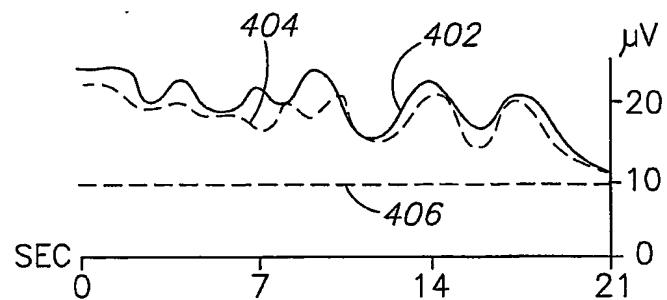


FIG.5A

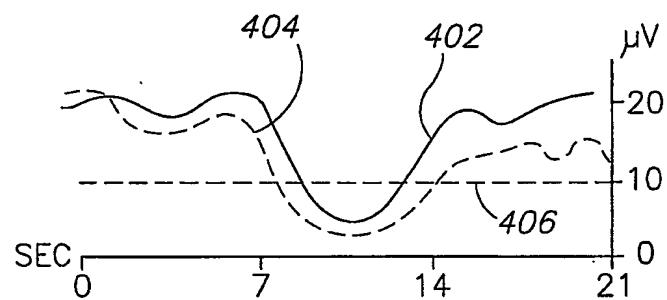


FIG.5B

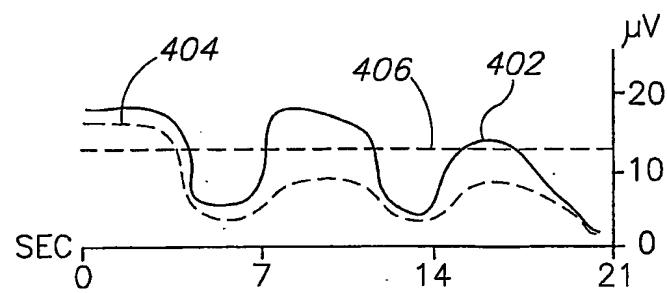


FIG.5C

5/5

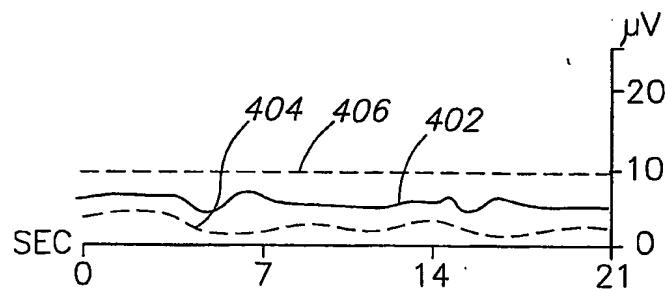


FIG.5D

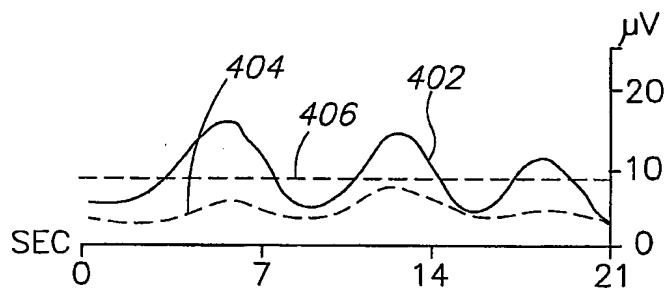


FIG.5E

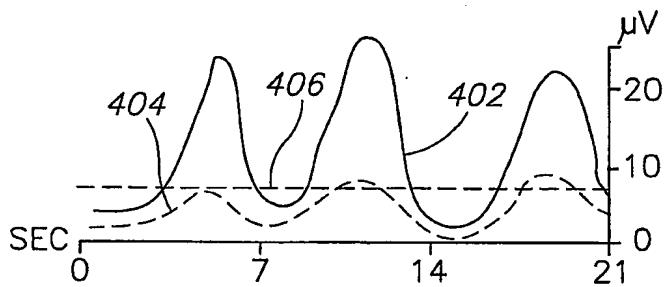


FIG.5F

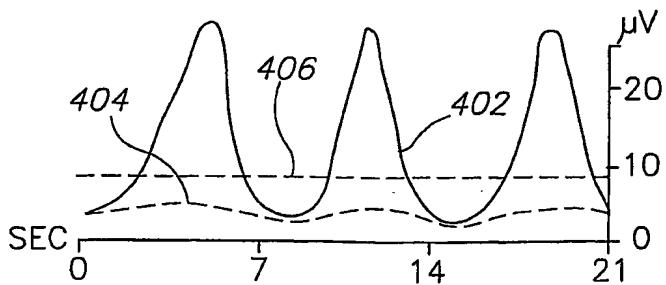


FIG.5G

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

**(19) World Intellectual Property Organization  
International Bureau**



A standard linear barcode is located at the bottom of the page, spanning most of the width. It is used for document tracking and identification.

**(43) International Publication Date  
22 September 2005 (22.09.2005)**

PCT

(10) International Publication Number  
**WO 2005/087307 A3**

**(51) International Patent Classification:**  
*A61N 1/36 (2006.01) A61B 5/0488 (2006.01)*

(81) **Designated States** (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TI, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.

**(21) International Application Number:** PCT/IL2005/000135

(84) **Designated States** (*unless otherwise indicated, for every kind of regional protection available*): ARIPO (BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

(22) International Filing Date: 4 February 2005 (04.02.2005)

**(25) Filing Language:** English

**(26) Publication Language:** English

**(30) Priority Data:**

60/542,022 5 February 2004 (05.02.2004) US  
60/566,078 29 April 2004 (29.04.2004) US

(71) **Applicant (for all designated States except US):** MO-TORIKA INC. [GB/GB]; Nerine Chambers, P.O. Box 905, Road Town, Tortola (VG).

**(72) Inventors; and**

(75) **Inventors/Applicants (for US only):** EINAV, Omer [IL/IL]; 42875 Kfar Monash (IL). KORENMAN, Ernesto [IL/IL]; 7 Hativat Givati Street, 43338 Raanana (IL).

(74) **Agents:** FENSTER, Paul et al.; FENSTER & COMPANY, INTELLECTUAL PROPERTY LTD., P.O. Box 10256, 49002 Petach Tikva (IL).

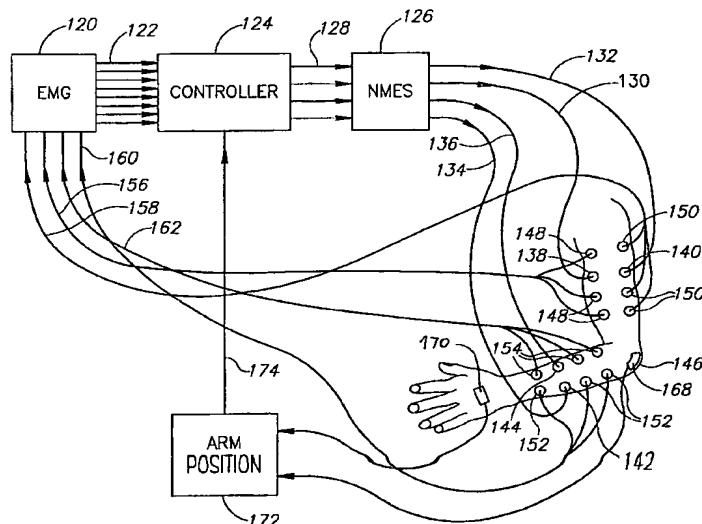
**Published:**

— *with international search report*

**(88) Date of publication of the international search report:**  
3 May 2007

*For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.*

**(54) Title: NEUROMUSCULAR STIMULATION**



**(57) Abstract:** Apparatus for rehabilitating a patient who has a paretic body part (146), the apparatus comprising: a) at least one electromyography (EMG) sensor adapted, to being applied to a voluntary muscle of a healthy body part of the same type as the paretic body part, which at least one sensor produces at least one EMG signal; b) a neuromuscular electrical stimulation (NMES) device (126) adapted for stimulating at least one voluntary muscle of the paretic body part; and c) a controller (124) which controls the NMES device, making the amplitude of stimulation of the paretic body part at least partly dependent on the EMG signal from the healthy body part. In another embodiment, where only a paretic body part is used, the amplitude of stimulation is not sufficient to cause contraction by itself but only together with the patient's own nerve impulses.

WO 2005/087307 A3

**INTERNATIONAL SEARCH REPORT**

International Application No

PCT/IL2005/000135

**A. CLASSIFICATION OF SUBJECT MATTER**  
IPC 7 A61N1/36 A61B5/0488

According to International Patent Classification (IPC) or to both national classification and IPO

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61N A61B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, PAJ

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category <sup>a</sup>	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 02/092164 A (FONDATION SUISSE POUR LES CYBERTHESES; BODDARD, ROLAND; CLAVEL, REYMON) 21 November 2002 (2002-11-21) page 14, paragraph 8 - page 15, paragraph 3 page 17, paragraph 7 - page 18, paragraph 2 page 18, paragraph 5 page 19, paragraph 4 - page 20, paragraph 5; figure 1 ----- -/-	1-16, 18, 19, 23-30
A	-----	17, 20-22, 31

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

<sup>a</sup> Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the International filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
- "&" document member of the same patent family

Date of the actual completion of the International search

23 May 2005

Date of mailing of the International search report

03/06/2005

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2  
NL - 2280 HV Rijswijk  
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,  
Fax: (+31-70) 340-3016

Authorized officer

Küster, G

## INTERNATIONAL SEARCH REPORT

International Application No  
PCT/IL2005/000135

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 4 582 049 A (YLVISAKER ET AL) 15 April 1986 (1986-04-15)	1-3, 8-10, 14-17
A	column 2, lines 25-35  column 2, line 51 - column 3, line 26 column 4, lines 17-43 column 5, lines 33-50; figure 1 -----	4-7, 11-13, 18-31
X	GRAUPE D: "EMG PATTERN ANALYSIS FOR PATIENT-RESPONSIVE CONTROL OF FES IN PARAPLEGICS FOR WALKER-SUPPORTED WALKING" IEEE TRANSACTIONS ON BIOMEDICAL ENGINEERING, IEEE INC. NEW YORK, US, vol. 36, no. 7, 1 July 1989 (1989-07-01), pages 711-719, XP000037463 ISSN: 0018-9294	1,3, 8-10,14, 16,17,19
A	page 711, left-hand column, paragraph 1 - right-hand column, paragraph 1; figures 3,5 page 716, left-hand column, paragraph 1 -----	31
X	WO 02/13673 A (STIMEL LTD; KOTLIK, BEN-ZION; ZUKER, MORRIS) 21 February 2002 (2002-02-21)	31-35
A	page 10, lines 25-32 page 11, line 30 - page 12, line 11 page 12, line 25 - page 13, line 1; figure 1 -----	1
X	PEASGOOD W ET AL: "EMG-controlled closed loop electrical stimulation using a digital signal processor" ELECTRONICS LETTERS, IEE STEVENAGE, GB, vol. 36, no. 22, 26 October 2000 (2000-10-26), pages 1832-1833, XP006015871 ISSN: 0013-5194	31-35
A	page 1832, left-hand column, paragraph 1; figure 1 page 1833, right-hand column, paragraph 1 -----	1

## INTERNATIONAL SEARCH REPORT

International application No.  
PCT/IL2005/000135

### Box II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1.  Claims Nos.: **36-46** because they relate to subject matter not required to be searched by this Authority, namely:  
**Rule 39.1(iv) PCT - Method for treatment of the human or animal body by therapy**
2.  Claims Nos.: because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3.  Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

### Box III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1.  As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2.  As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3.  As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4.  No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

#### Remark on Protest

The additional search fees were accompanied by the applicant's protest.  
 No protest accompanied the payment of additional search fees.

**INTERNATIONAL SEARCH REPORT**

Information on patent family members

International Application No

PCT/IL2005/000135

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
WO 02092164	A 21-11-2002	CA 2446875	A1 21-11-2002	
		WO 02092164	A2 21-11-2002	
		EP 1387712	A2 11-02-2004	
		JP 2004526541	T 02-09-2004	
		US 2004172097	A1 02-09-2004	
<hr/>				
US 4582049	A 15-04-1986	DE 3516279	A1 13-11-1986	
		FR 2580934	A1 31-10-1986	
		GB 2175504	A 03-12-1986	
<hr/>				
WO 0213673	A 21-02-2002	AU 8007001	A 25-02-2002	
		EP 1311320	A2 21-05-2003	
		WO 0213673	A2 21-02-2002	
		US 2003208246	A1 06-11-2003	
<hr/>				